We claim:

1 1. A pharmaceutical composition for the treatment of a bacterial infection in a
2 mammal which comprises a therapeutically effective amount of a compound having the formula

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wherein:

R₁ is hydrogen, alkyl, alkanoyl or Y-substituted alkanoyl

wherein Y is alkyl, aryl or halo; and

 R_2 is amide, or X-substituted amide wherein X is a peptide or an amino acid; or a pharmaceutically acceptable addition salt and/or hydrate thereof, or where applicable, a geometric or optical isomer or racemic mixture thereof.

- 2. The pharmaceutical composition of Claim 1 wherein R_1 is alkanoyl and R_2 is X-substituted amide wherein X is an amino acid residue.
 - 3. The pharmaceutical composition of Claim 1 wherein R_1 is acetyl and R_2 is prolyl.
- 4. The pharmaceutical composition of Claim 1 wherein said compound has the
 formula: 3β-acetoxy-17β-(L-prolyl)amino-5α-androstane.
- 1 5. The pharmaceutical composition of Claim 1 and a pharmaceutically acceptable 2 carrier.
- 1 6. A pharmaceutical composition according to claim 5 in a form suitable for topical administration.

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- 1 7. A pharmaceutical composition according to Claim 5 wherein said carrier is 2 selected from the group comprising lotion, salve, ointment, cream or oil.
- 1 8. A pharmaceutical composition according to Claim 1 comprising in addition a 2 second anti-microbial agent.
- 9. A pharmaceutical composition according to Claim 5 comprising in addition
 means for controlling the pH of said composition.
- 1 10. A method of treating a gram positive bacterial infection in a mammal which 2 comprises administering to said mammal an antimicrobial-effective amount of a compound of 3 claim 5.
 - 11. The method of Claim 10 wherein said antimicrobial-effective amount is between about 25 milligram to about 1 gram per kilogram body weight of said mammal treated.
 - 12. A method of inhibiting the growth of gram positive bacteria comprising contacting said bacteria with a compound of Claim 1.
 - 13. The method of Claim 10 wherein said gram-positive bacteria are selected from the group comprising penicillin-resistant, methicillin-resistant and vancomycin resistant gram-positive bacteria.
- 1 14. The method of Claim 10 wherein said compound is administered to said mammal 2 by topical means.
- 1 15. The method of Claim 14 wherein said means is selected from the group comprising lotion, oil, emulsion and creme.
- 1 16. The method of Claim 14 wherein said means comprises a surface-adhering 2 dressing impregnated with said compound.